

Virtual Testing of Concrete Transport Properties

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Synopsis: The transport properties of concrete are critical to its field performance. Commonly encountered degradation mechanisms are dependent on ionic diffusivity, sorptivity, and permeability. In this paper, virtual testing of two of these concrete transport properties, diffusivity and permeability, will be reviewed. Virtual evaluations of ionic diffusion (and equivalently conductivity) will be presented as one example that spans the full range of applications, from computations on cement paste with micrometer resolution to a virtual rapid chloride permeability test (RCPT) that simulates the standard ASTM test method for conductivity of concrete cylinders. At the concrete scale, a hard core/soft shell (HCSS) microstructural model may be employed to estimate diffusion coefficients, while finite difference solutions of Fick’s laws that incorporate sorption/reaction may be employed to evaluate remediation strategies for real world bridge decks. Virtual evaluations of permeability are dependent on a sufficient resolution of the pore sizes that are critical for flow under pressure. Two recent successful evaluations will be presented in this paper: the permeability of cement pastes (hydroceramics) cured at elevated temperatures, where transport is controlled by micrometer-sized pores, and the permeability of pervious concrete that is dominated by its coarse porosity (scale of mm). Many of the presented computational (virtual) tools are freely available over the Internet, either for direct access (remote computation) or for downloading.

Keywords: Building technology; diffusion; durability; microstructure; permeability; transport; virtual testing.

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INTRODUCTION

To engineer durable concrete structures that will provide an acceptable service life, appropriate mechanical and transport properties are equally critical.¹ All common concrete degradation processes including corrosion of steel reinforcement, sulfate attack, alkali-silica reaction, and damage due to freezing/thawing (scaling) are strongly influenced by one or more of the following three major transport mechanisms: diffusion, permeability, and sorptivity. Thus, a viable approach to increase concrete service life is to reduce the values of these transport coefficients. Understanding the relationships between microstructure and transport can accelerate the achievement of such reductions. In this paper, approaches for obtaining an increased understanding of diffusion and permeability of cement-based materials based on computational materials science will be presented. This virtual testing of concrete properties can provide deeper insights into microstructure-property relationships and can also reduce expensive, laborious experimental testing.²

RESEARCH SIGNIFICANCE

While computational materials science has been applied to cement-based materials for well over twenty years, it generally has focused more on fundamental scientific issues rather than real world practical scenarios. This paper will present results that bridge this gap between computational materials science and real world applications. In addition to having immediate application to the real world problems, these results will demonstrate the powerful (and real) potential of a computational materials-science-based approach and hopefully aid in demystifying the modeling of concrete materials and performance.

MODELING TECHNIQUES

Microstructural Models

While the focus of this paper is on transport, a brief introduction into three different microstructural models that will be employed to provide inputs to algorithms for computing transport will be presented. The first, CEMHYD3D, is a computer model for the hydration and microstructure development of cement paste that is founded on a digital-image-based approach in which a three-dimensional computational volume is divided up into a cubic grid of voxels, where each voxel contains a single cementitious phase (starting material, water, or hydration product).^{3,4} The hydration process is simulated by iteratively applying a set of rules to mimic dissolution, diffusion, and precipitation throughout the three-dimensional set of voxels. A typical CEMHYD3D microstructure is 100 elements on a side (or 1 million voxels in total) with a voxel dimension of 1 μm (3.94×10^{-5} in). Because the model microstructure is available as a three-dimensional digital image, it can be easily utilized as input to finite element or finite difference algorithms for computing mechanical or transport properties. At this voxel size, the CEMHYD3D model exhibits a percolation/depercolation transition for the capillary porosity at a value of about 20 %, in general agreement with experimental data for ordinary portland cement pastes.⁵ In a percolated system, there is a continuous pathway across the microstructure within the capillary porosity. In a depercolated system, there is no such pathway within the porosity from one side of the microstructure to the opposite one.

The second microstructural model is the so-called hard core/soft shell (HCSS) model, one documented version of which is readily available from NIST.⁶ As opposed to CEMHYD3D, HCSSModel is a continuum-based model consisting of hard impenetrable (spherical) particles, optionally surrounded by a soft shell. Originally developed to represent concrete aggregates and their surrounding interfacial transition zones (ITZ),⁶ the model has been extended and adapted for numerous concrete applications including the mitigation of spalling of high-performance concretes via the addition of polymeric fibers,⁷ the adaptation of the protected paste volume concept to internal curing,⁸ and most recently the documentation of the influence of water-cement ratio (w/c) and cement particle size distribution on particle spacing in fresh cement paste.⁹ The model represents a three-dimensional cube of concrete (mortar or paste) and the particle size distribution for the particles is chosen to match that of the real materials. For example, models as large as a 50 mm x 50 mm x 50 mm (2 in x 2 in x 2 in) cube of concrete have been created, requiring well over 1 million individual particles to adequately represent a realistic concrete aggregate size distribution. For the HCSSModel, the percolation/depercolation transition for the “bulk” phase (not contained in the hard cores or the soft shells) occurs at a volume fraction of 3 % to 4 %, as it is more difficult to disconnect the bulk phase by concentric spherical shells than it is by the random growth processes employed in the CEMHYD3D model.^{10,11}

The third microstructural model is a general image-based reconstruction algorithm for creating a three-dimensional microstructural representation based on a two-dimensional image, as obtained using scanning electron microscopy, for example.¹² Assuming an isotropic material, the autocorrelation function measured on the porosity phase in the two-dimensional image is utilized to reconstruct a “representative” three-dimensional microstructure with the same porosity, surface area, and correlation as the original material. While first developed by Joshi¹³ and then extended by Quiblier¹⁴ many years ago, very recently these algorithms have found renewed applications to cement-based materials, including pervious concrete¹¹ and hydroceramics¹⁵ (high temperature cements for oil and geothermal well applications) as will be demonstrated in the results to follow. For the reconstruction-based model, the percolation threshold for the “porosity” phase has been observed to be on the order of 10 %, a value intermediate between the CEMHYD3D and HCSSModel values.¹²

Diffusivity/Conductivity

Based on the Nernst-Einstein equation shown in equation (1), diffusion and conduction in porous microstructures containing a conductive solution can be solved using the same basic computational procedures.¹⁶

$$\frac{\sigma}{\sigma_0} = \frac{D}{D_0} \quad (1)$$

where σ is the conductivity of the microstructure, σ_0 is the conductivity of the pore solution, D is the diffusivity of a specific ion in the microstructure, and D_0 is the corresponding diffusion coefficient for the ion in bulk solution of identical composition. Based on this equation, an electrical analogy can be used to estimate the diffusivity of a three-dimensional cement paste microstructure such as that produced by CEMHYD3D, as shown in the top portion of Figure 1. Adjacent voxels in the microstructure are simply connected together by resistors, where the resistance level indicates the relative conductivity (diffusivity) of the phases of the two voxels being connected. Specifically, resistors connecting together two capillary pore voxels are assigned a value of 1, while those connecting two voxels of the calcium silicate hydrate gel (C-S-H) are typically assigned a value of 250 to 400.^{16,17} The resultant electrical network can then be solved by finite element or finite difference techniques to obtain the conductivity (diffusivity) of the complete microstructure.¹⁸ Such an approach was used in the past to develop an equation for the diffusivity of tricalcium silicate pastes as a function of capillary porosity,¹⁶ an approach that was later extended to cement pastes, including those with silica fume additions.¹⁷

For a continuum-based microstructure, such as one generated by the HCSSModel, applying the electrical analogy is difficult. Instead, a continuum random walker (blind ant) approach is commonly employed for estimating diffusivities (conductivities) as indicated in the middle portion of Figure 1.^{6,19,20} In these simulations, each blind ant carries his own watch and global positioning system (GPS) and steps randomly (and blindly) through the microstructure, with steps in a more porous ITZ taking less time than equivalent steps in the bulk cement paste, for example. After a large number (hundreds of thousands to millions) of steps, the diffusivity of the microstructure can be estimated from the mean distance squared per unit time achieved by all of the ants (Figure 1). For a typical simulation, 10 000 ants are used. The HCSSModel provides an extremely flexible format for simulating real world variations, such as a concrete with internal curing, where the lightweight aggregates are not surrounded by the ITZs that are present around their normal weight counterparts, producing a significant reduction in the diffusion of chloride ions.²¹

Based on computations performed on virtual cement paste and concrete,^{17,20} the next logical step is to proceed to virtual test procedures, such as simulating the response of a field concrete to a standard ASTM or American Association of State Highway and Transportation Officials (AASHTO) rapid chloride permeability test (RCPT) as indicated in the bottom portion of Figure 1.²² Such a virtual test has been developed by NIST and is available at <http://ciks.cbt.nist.gov/VirtualRCPT.html>. The interactive application allows the user to select their mixture proportions and testing conditions and provides predictions of the pore solution composition, its conductivity,²³ and ultimately the RCPT response. Features include the ability to reduce the testing time from the standard 6 h to just a few minutes, to avoid temperature effects, as suggested by Snyder et al.²⁴ in 2000 and more recently advocated by Riding et al.²⁵ An example of the predictive ability of the virtual RCPT test is provided in Figure 2, which contrasts the experimental and virtual RCPT values²² for a variety of concretes from one particular study.²⁶

The final step in the modeling of diffusion is the prediction of chloride ion profiles obtained under realistic field exposure conditions. NIST has developed a web-based application (Computer Integrated Knowledge System-CIKS) for providing such predictions (<http://ciks.cbt.nist.gov/~bentz/millandfill/clpenmillandfill.html>).²⁷ The model is based on a one-dimensional finite difference solution for diffusion that includes chloride ion binding by the cementitious hydration products and reaction with the aluminate phases to form Friedel's salt. The user specifies the average exposure temperature and chloride loading for each month of the year, as well as a variety of mixture proportioning parameters.²⁷ Working with Brigham Young University, NIST has recently extended this application to consider real world remediation strategies such as surface treatments²⁸ and scarification and overlays (SO),²⁹ as will be presented further in the results section to follow.

Permeability

The basic goal in computing the permeability of a material is to apply a pressure gradient across the three-dimensional microstructure and compute the resultant flow. NIST has developed and recently distributed a three-dimensional linear Stokes solver for performing this calculation for a three-dimensional microstructure consisting of pores and solids.³⁰ A finite difference solution of the linear Stokes equations for slow, incompressible, steady-state flow is utilized to determine the x, y, and z components of the fluid velocity in each porosity voxel. Once this finite difference solution converges sufficiently, the intrinsic permeability, k , of the composite three-dimensional microstructure is calculated by volume averaging the local fluid velocity (in the direction of the flow) and applying the Darcy equation:

$$u = -\frac{k}{\eta} \frac{\Delta P}{L} \quad (2)$$

where u is the average fluid velocity in the direction of the flow (x-direction) for the microstructure obtained via the Stokes solver program, η is the fluid viscosity, and L is the length of the sample microstructure across which an applied pressure difference of ΔP exists.³¹ For a given microstructure, three separate runs of the computer codes may be conducted to determine the (different) permeabilities for pressure-driven flow in the x, y, and z directions. Obviously, for permeability predictions to be accurate, the three-dimensional microstructure used as input must adequately represent the pores (sizes, connectivity, and tortuosity) that are dominating the flow paths through the material.

RESULTS AND DISCUSSION

Diffusivity/Conductivity

Many state departments of transportation (DOTs) are concerned with the penetration of chlorides (from de-icing salts, for example) into bridge deck concretes and the resultant corrosion problems and concrete degradation. Two approaches that are often considered for remediation of existing bridge decks are the application of impermeable surface treatments to prevent ingress of future chlorides and the scarification (milling) of the existing bridge deck followed by the application of an overlay (filling), to remove the most chloride-laden portion of the concrete and replace it by a (thicker) high-performance concrete.^{28,29} As mentioned above, the CIKS-Chloride computer model has been recently modified to simulate the effects of both of these remediation strategies.

The application of these computer models has been demonstrated in a project that considered differences in predicted performance for various cover depths for bridge decks with and without stay-in-place metal forms (SIPMFs).^{28,29} Relevant chloride ion diffusion coefficients were determined from chloride-ion profile measurements obtained on specimens from 12 existing concrete bridge decks (six without SIPMFs and six with SIPMFs). The CIKS system was then employed to simulate chloride concentrations at the (variable) depth of cover as a function of the exposure age and the time of application of a given remediation strategy. Example results are shown in Figures 3 and 4 for the case of a 50 mm (2 inch) concrete cover. When SIPMFs are employed, higher diffusion coefficients are measured, due to the increased saturation level of the concrete. This requires the remediation strategies to be employed at earlier times to provide the same performance. Table 1 summarizes the latest time at which successful remediation can be achieved with the performance criteria that the chloride concentration at the level of the top layer of reinforcement (cover depth) does not exceed 1.18 kg/m³ (2 lb/yd³). Because the scarification and overlay treatment actually removes a portion of the chlorides (in the removed top layer of the concrete), treatment can be slightly delayed when this more severe (and expensive) remediation strategy is employed. However, the differences between treatments are far outweighed by the differences in deck type due to the increase in chloride ion diffusion coefficients for decks with SIPMFs compared to those without SIPMF. This study thus emphasizes the criticality of producing a concrete with a low chloride ion diffusivity.

Permeability

Originally, the permeability computation codes were developed with the hopes of computing permeabilities as a function of degree of hydration for cement paste microstructures. While the predictions for fresh cement pastes were reasonable, experimental permeabilities fell dramatically as the capillary pores depercolated during hydration and the digital-image-based cement paste microstructural model was not able to adequately resolve a wide enough

range of pores sizes to provide reasonable permeability predictions. The permeability codes then remained dormant for a number of years until recently when two new applications were found: cement pastes cured at elevated temperatures (hydroceramics) and pervious concrete.

In deep oil and geothermal well cementing applications, hydration occurs at vastly elevated temperatures, as the wells have typical working temperatures in the range of 200 °C (392 °F) to 350 °C (662 °F).¹⁵ This requires the utilization of special cements and leads to the formation of a variety of hydration products not typically encountered for room temperature hydration.¹⁵ In addition, the porosities of these materials are generally higher and the pore sizes larger than those in a well hydrated ordinary portland cement. Thus, application of the permeability computation procedures to these hydroceramic materials was of interest. The general procedure employed in reference 15 is summarized in Figure 5. Starting from a two-dimensional scanning electron microscopy (SEM) image, a binary (pore/solid) image is obtained and analyzed by determining the two-dimensional correlation function for the porosity phase. This correlation function and the experimentally measured porosity for the material (values in the range of 25 % to 40 %) are then used to reconstruct a three-dimensional representation of the microstructure based on the procedures described above. The permeability computation code is then employed to compute a fluid permeability for the reconstructed microstructure.

Experimental and computational results are compared in Figure 6 for hydroceramics cured at 200 °C (392 °F) and containing either alumina or silica additions. While the alumina creates a more porous and permeable microstructure, the silica produces a denser microstructure with a dramatically reduced permeability. For both silica and alumina additions, the experimental trends are well captured by the estimates provided by the computational procedures. Further, the predictions are observed to be reasonably accurate for permeabilities spanning four orders of magnitude.

A second application where permeability is controlled by pores in a very limited size range is that of pervious concrete. Pervious concrete is basically produced by removing the fine aggregate fraction from a concrete mixture, producing a concrete with enhanced porosity and a set of connected pores that are up to several millimeters in size. In this case, an optical microscope can be utilized to capture a two-dimensional image of the porosity of the pervious concrete. The same two-dimensional image to three-dimensional microstructure algorithms can then be employed as shown in Figure 5. Figure 7 provides an overview of the process for a pervious concrete, along with a graph comparing previously obtained experimental and model permeability values. Once again, the predictions provided by the reconstruction/computation algorithms are in reasonable agreement with the experimental values compiled from three different sources.³²⁻³⁴ Conversely, as shown in the graph in Figure 7, model predictions generated based on a version of the HCSSModel produced permeabilities that were well above those measured experimentally, most likely due to that models much lower percolation threshold as mentioned earlier. In addition to permeability, the reconstructed three-dimensional microstructures of the pervious concrete can also be analyzed to assess other issues such as the potentials for clogging and freezing/thawing damage.¹¹

SUMMARY AND PROSPECTUS

Virtual testing of concrete transport properties aids in the development of fundamental microstructure/property relationships, such as those between transport coefficients and porosity (Figure 7) or other microstructural characteristics. Additionally, virtual testing has the potential to be applied to a wide variety of “what if” field scenarios to optimize mitigation and remediation strategies, as demonstrated for remediation of chloride-contaminated bridge deck concretes (Figures 3 and 4). The various models described in this paper and their locations on the Internet are summarized in Table 2. All are freely available for execution and/or downloading; documentation in the form of user manuals or accompanying technical articles is also provided.

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Table 1- Latest treatment times vs. bridge deck type and remediation strategy

Bridge deck type	Treatment type	Latest treatment time
SIPMF	Surface	1 year
No SIPMF	Surface	5 years
SIPMF	Scarification and overlay	2 years
No SIPMF	Scarification and overlay	6 years

Table 2- NIST models for virtual testing (available for remote execution or downloading)

Model	Description/Purpose	Download location or web page address
CEMHYD3D	Cement hydration (version 3.0)	ftp://ftp.nist.gov/pub/bfrl/bentz/CEMHYD3D/version30
HCSSModel	Hard core/soft shell 3-D microstructural model	http://ciks.cbt.nist.gov/emml.html
3-D Reconstruction	Correlation-based transformation from 2-D image to 3-D microstructure	ftp://ftp.nist.gov/pub/bfrl/bentz/permsolver/
VirtualRCPT	Virtual RCPT test	http://ciks.cbt.nist.gov/VirtualRCPT.html
CIKS-Chloride	Chloride penetration simulation including surface treatment and/or scarification and overlay	http://ciks.cbt.nist.gov/~bentz/millandfill/clpenmillandfill.html
Permsolver	3-D Stokes solver for permeability	ftp://ftp.nist.gov/pub/bfrl/bentz/permsolver/

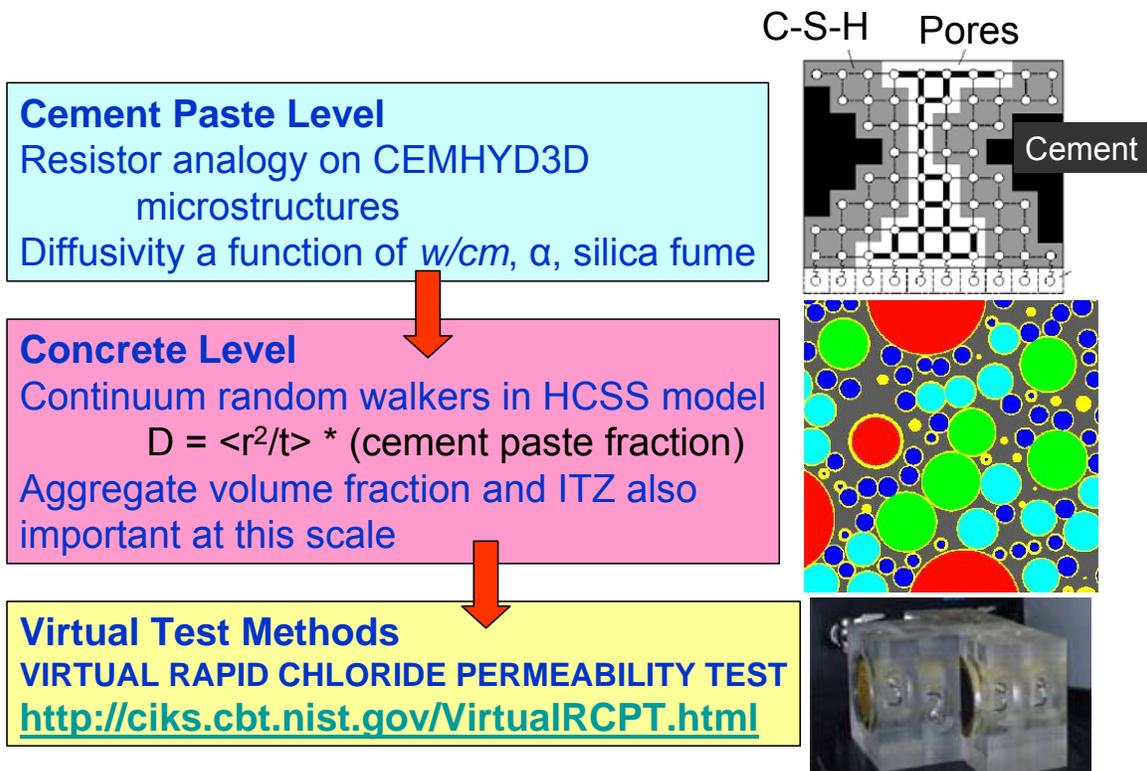


Fig. 1. Summary of multi-scale modeling of conductivity/diffusivity in cement-based materials.

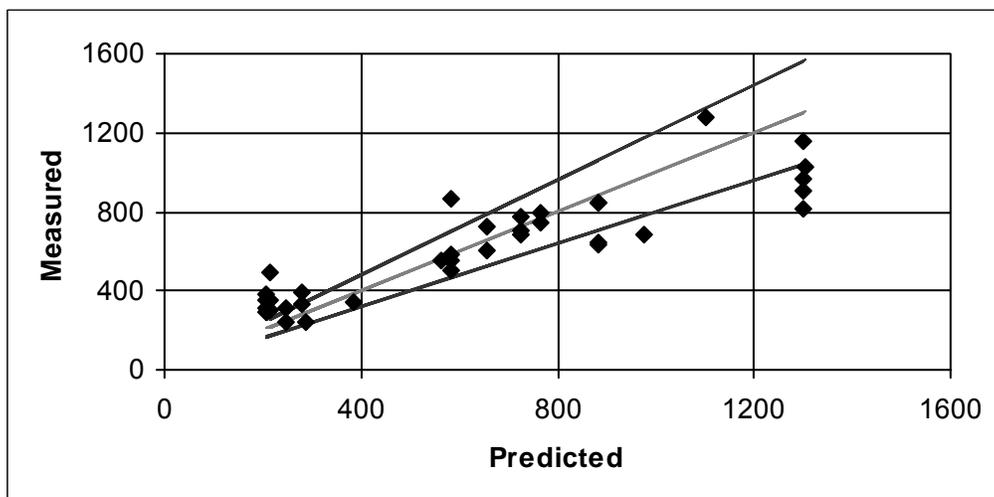


Fig. 2. Measured vs. predicted (virtual) RCPT test results.²² Experimental data taken from ref. 26. Grey line indicates line of equality, and dark lines indicate $\pm 20\%$ of the measured values.

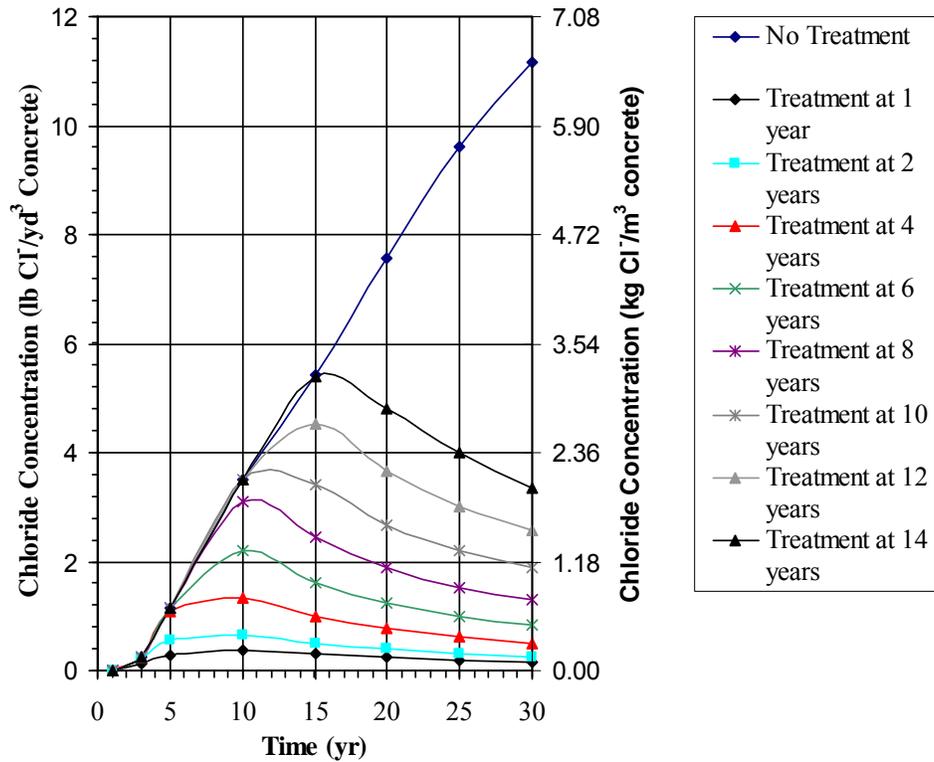
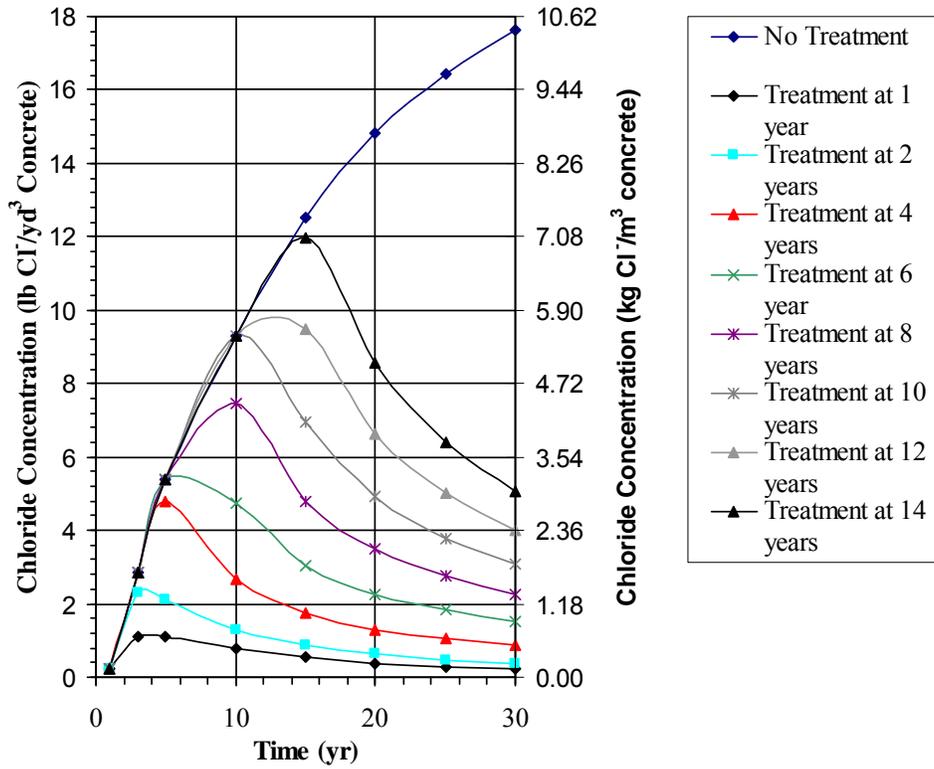


Fig. 3. Comparison of simulated chloride concentrations of decks with (top) and without (bottom) SIPMFs at 50 mm (2 in) cover depth as a function of time of application of surface treatment.²⁸ One kg/m³ = 1.69 lb/yd³.

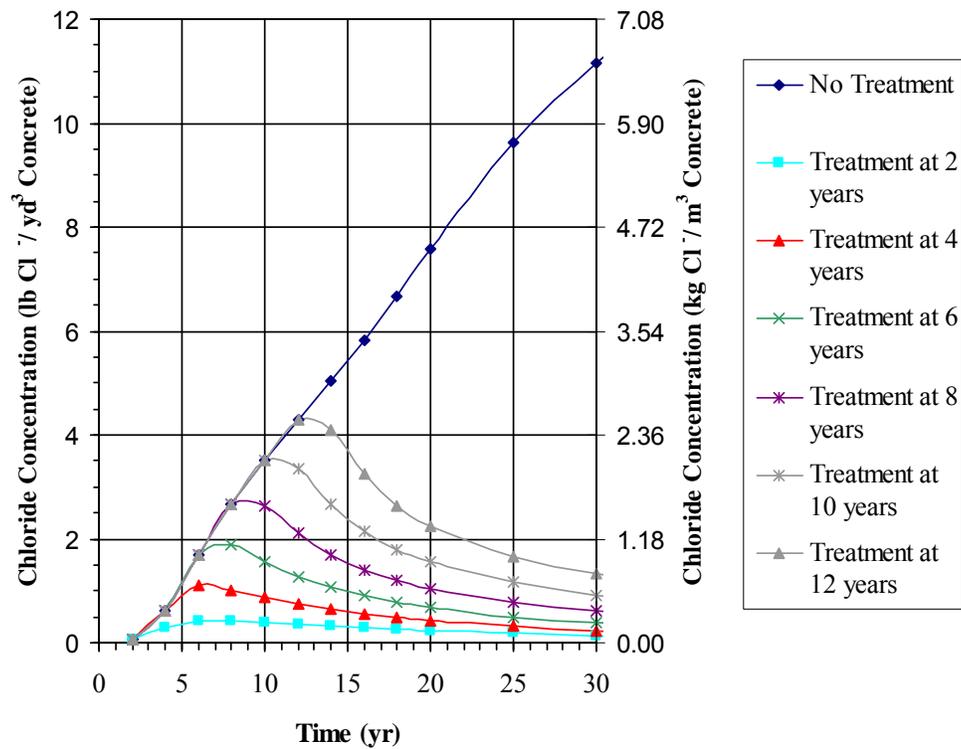
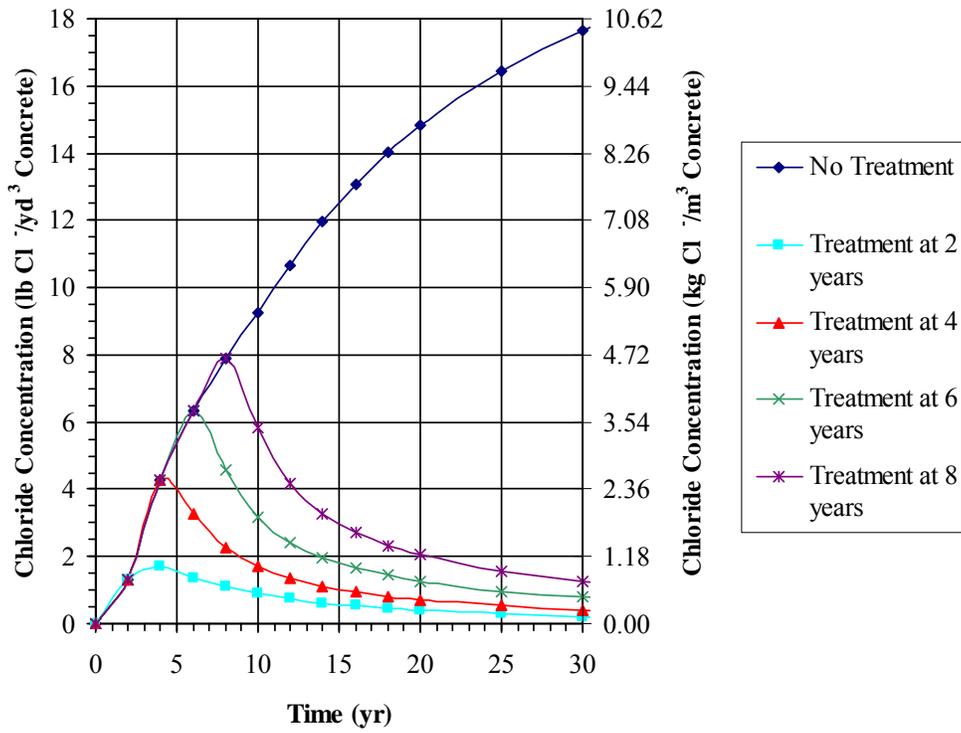


Fig. 4. Comparison of simulated chloride concentrations of decks with (top) and without (bottom) SIPMFs at 50 mm (2 inch) cover depth as a function of timing of a 12.7 mm (0.5 inch) scarification and 38 mm (1.5 inch) overlay treatment.²⁹ One kg/m³ = 1.69 lb/yd³.

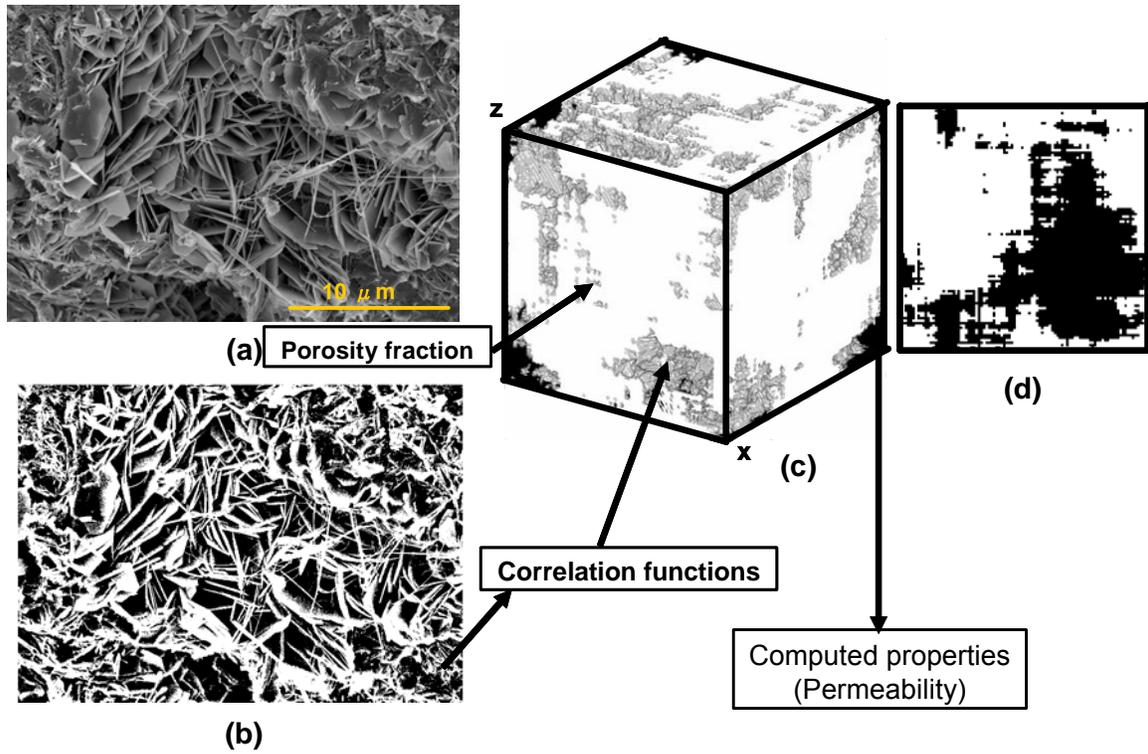


Fig. 5. Illustration of the process of creating a three-dimensional microstructure for subsequent computation of permeability.¹⁵ **a)** SEM image showing hexagonal crystals of truscottite. The needle shaped crystals are of xonotlite. Sample cured at 350 °C containing 40 % by mass of silica flour. **b)** Binary image used to extract the correlation functions for the 3-D reconstruction. **c)** Reconstructed three-dimensional microstructure (100 pixels x 100 pixels x 100 pixels) for which permeability is computed. **d)** Slice of the reconstructed three-dimensional microstructure.

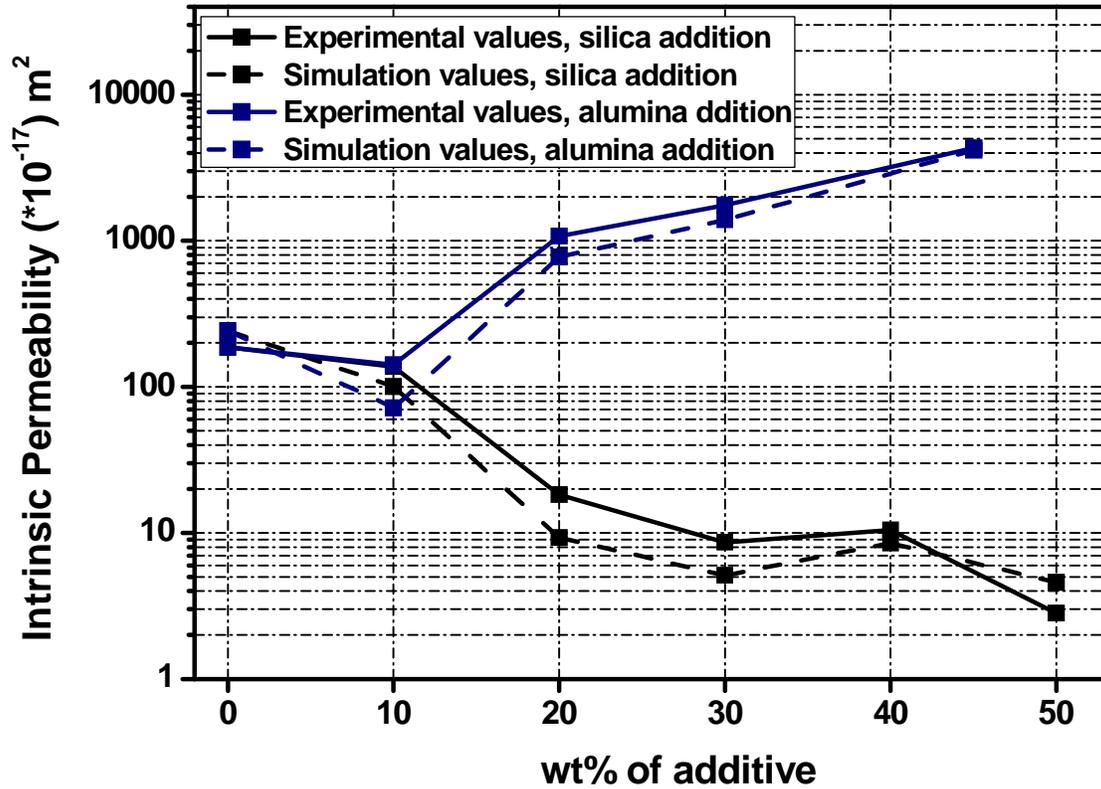


Fig. 6. Comparison of experimental and simulation values of intrinsic permeability with silica or alumina additions in hydroceramic samples.¹⁵ $1 \text{ m}^2 = 10.76 \text{ ft}^2$.

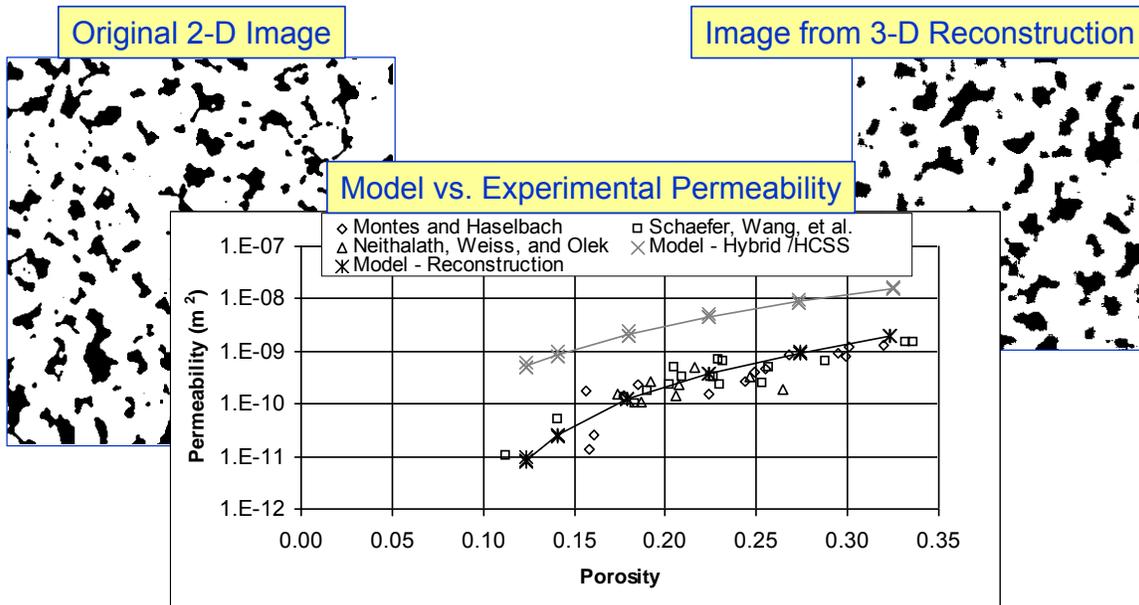


Fig. 7. Comparison of experimental and simulation values for intrinsic permeability for pervious concretes.¹¹ Original image is 63 mm by 63 mm (2.5 in by 2.5 in). Experimental values were extracted from the references listed in the legend.³²⁻³⁴ $1 \text{ m}^2 = 10.76 \text{ ft}^2$.